

**NIGHTTIME SLEEP AID- diphenhydramine hcl solution**  
**L.N.K. International, Inc.**

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**Quality Plus 44-002A**

***Active ingredient (in each 30 mL)***

Diphenhydramine HCl 50 mg

***Purpose***

Nighttime sleep-aid

***Uses***

- for relief of occasional sleeplessness
- reduces time to fall asleep if you have difficulty falling asleep

***Warnings***

**Do not use**

- for children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin

**Ask a doctor before use if you have**

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland

**Ask a doctor or pharmacist before use if you are**

taking sedatives or tranquilizers.

**When using this product**

avoid alcoholic beverages.

**Stop use and ask a doctor if**

sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.

**If pregnant or breast-feeding,**

ask a health professional before use.

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away.

***Directions***

- **do not take more than directed**
- mL = milliliter
- only use the dose cup provided
- take only one dose per day (24 hours)
- adults and children 12 years and over: take 30 mL at bedtime if needed or as directed by a doctor
- children under 12 years: do not use

***Other information***

- **each 30 mL contains:** sodium 17 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

***Inactive ingredients***

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavor, glycerin, high fructose corn syrup, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sucralose, sucrose, xanthan gum

***Questions or comments?***

**1-800-426-9391**

***Principal display panel***

**Quality**

**+ Plus**

NDC 50844-200-02

\*Compare to active ingredient in VICKS® ZzzQuil® NIGHTTIME SLEEP-AID

**Nighttime**

**Sleep Aid**

**Diphenhydramine HCl 50 mg per 30 mL**

Nighttime Sleep-Aid

- Non-habit Forming
- Not For Colds or For Pain

Berry Flavor

Ages 12 Years and Over

12 FL OZ (355 mL)

**TAMPER EVIDENT: DO NOT USE IF PRINTED  
NECK WRAP IS BROKEN OR MISSING**

\*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark VICKS® ZzzQuil® NIGHTTIME SLEEP-AID. 50844 ORG022300202

Distributed by  
**LNK INTERNATIONAL, INC.**  
60 Arkay Drive  
Hauppauge, NY 11788  
USA

**QUALITY PLUS** NDC 50844-200-02  
\*Compare to active ingredient in VICKS® ZzzQuil® NIGHTTIME SLEEP-AID

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F-002A ORG

PEEL BACK FOR COMPLETE DRUG FACTS

**Drug Facts** TAMPER EVIDENT: DO NOT USE IF PRINTED  
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B-002A ORG 3 50844 20002 0

no print / no varnish area  
lot no. & exp. date

**Drug Facts** (continued)

When using this product avoid alcoholic beverages.

Stop use and ask a doctor if sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.

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**Questions or comments?**  
1-800-426-9391

Quality Plus 44-002A

**NIGHTTIME SLEEP AID**

diphenhydramine hcl solution

**Product Information**

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:50844-200

<b>Route of Administration</b>	ORAL
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### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>HIGH FRUCTOSE CORN SYRUP</b> (UNII: XY6UN3QB6S)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

### Product Characteristics

<b>Color</b>	purple	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BERRY	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50844-200-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	03/31/2025
2	NDC:50844-200-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2016	
3	NDC:50844-200-96	2 in 1 PACKAGE	09/12/2016	
3		355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M010	09/12/2016	
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**Labeler -** L.N.K. International, Inc. (038154464)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LNK International, Inc.		967626305	manufacture(50844-200) , pack(50844-200)

Revised: 3/2025

L.N.K. International, Inc.